Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

- 1. (Currently Amended) A polyvalent multimeric complex comprising a plurality of chimeric pRNA monomers, each said chimeric pRNA monomer independently comprising a heterologous component that comprises a biologically active RNA.
- 2. (Currently Amended) The polyvalent multimeric complex of claim 1 wherein the heterologous component of at least one chimeric pRNA monomer comprises a biologically active RNA <u>is</u> selected from the group consisting of a ribozyme, a siRNA, an RNA aptamer, an antisense RNA and a peptide nucleic acid (PNA).
- 3. (Previously Presented) The polyvalent multimeric complex of claim 1 wherein the heterologous component of at least one chimeric pRNA monomer comprises an endlabeling agent.
- 4. (Previously Presented) The polyvalent multimeric complex of claim 3 wherein the end-labeling agent is selected from the group consisting of biotin, pCp, DIG, SH group and phosphate.
- 5. (Previously Presented) The polyvalent multimeric complex of claim 1 wherein at least one of the chimeric pRNA monomers is a circularly permuted pRNA.
 - 6. (Canceled)

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- 7. (Previously Presented) The polyvalent multimeric complex of claim 1 wherein at least one of the chimeric pRNA monomers incorporates at least one nucleotide analog or modified nucleotide.
- 8. (Previously Presented) The polyvalent multimeric complex of claim 7, wherein the nucleotide analog or modified nucleotide is selected from the group consisting of a 2'-F-2' deoxy nucleotide derivative, a phosphorothoiate, a 2'-O-methyl ribonucleotide, a peptide nucleic acid (PNA).
 - 9.-16. (Canceled)
- 17. (Withdrawn) A method for delivering a therapeutic agent to a cell comprising:

contacting the cell with the polyvalent multimeric complex of claim 1, wherein the heterologous component of a first chimeric pRNA monomer comprises a therapeutic agent and the heterologous component of a second chimeric pRNA monomer comprises a biologically active moiety that specifically binds a component of the cell membrane, such that the polyvalent multimeric complex is taken up by the host cell.

- 18. (Withdrawn) The method of claim 17 wherein the component of the cell membrane to which the polyvalent multimeric complex binds is a receptor, and wherein the polyvalent multimeric complex is taken up by the cell via receptor-mediated endocytosis.
 - 19.-27. (Canceled)
- 28. (Currently Amended) A non-circularly permuted chimeric pRNA monomer comprising 5' and 3' ends, wherein at least one of said 5' and 3' ends comprises a heterologous component that comprises a biologically active RNA, and wherein the chimeric pRNA monomer is capable of assembling into a polyvalent multimeric complex comprising a plurality of chimeric pRNA monomers.

- 29. (Previously Presented) The pRNA monomer of claim 28 wherein the heterologous component comprises antisense RNA.
- 30. (Previously Presented) The pRNA monomer of claim 28 wherein the heterologous component comprises an aptamer.
- 31. (Previously Presented) The pRNA monomer of claim 28 wherein the heterologous component comprises a labeling agent.
- 32. (Previously Presented) The pRNA monomer of claim 31 wherein the labeling agent is selected from the group consisting of biotin, pCp, DIG, SH group and phosphate.
- 33. (Previously Presented) The pRNA monomer of claim 32 comprising at least one nucleotide analog or modified nucleotide.
- 34. (Previously Presented) The pRNA monomer of claim 33, wherein the nucleotide analog or modified nucleotide is selected from the group consisting of a 2'-F-2' deoxy nucleotide derivative, a phosphorothioate, a 2'-O-methyl ribonucelotide, a peptide nucleic acid (PNA).
- 35. (Currently Amended) <u>A The chimeric pRNA monomer of claim 28</u> comprising at least one nucleotide analog or modified nucleotide.
- 36. (Previously Presented) The chimeric pRNA monomer of claim 35 which is a circularly permuted pRNA.
- 37. (Previously Presented) The chimeric pRNA monomer of claim 35 wherein the nucleotide analog or modified nucleotide is selected from the group consisting of a

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2'-F-2' deoxy nucleotide derivative, a phosphorothioate, a 2'-O-methyl ribonucleotide, a peptide nucleic acid (PNA).

38.-48. (Canceled)